Listing of Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1-38. (Canceled).

39. (Currently Amended) A method for treating an autoimmune disorder, comprising administering to a subject having an autoimmune disorder**, an effective amount of** a therapeutic composition comprising a pharmaceutically acceptable carrier and at least one ~~antibody selected from the group consisting of an~~ **non-blocking** anti-CD22 antibody ~~which targets an A, B, D, or E epitope of CD22, an anti-CD20 antibody, and an anti-CD19 antibody, wherein said at least one antibody is administered in an amount effective for inactivating or depleting B-cells in said subject~~.

40. (Previously Presented)The method of claim 39, wherein said therapeutic composition is administered parenterally in a dosage of from 20 to 2000 mg per dose.

41. (Currently Amended) The method of claim **39** [[40]], wherein said subject receives said antibody in repeated parenteral dosages.

42. (Previously Presented) The method of claim 39, wherein said antibody is selected from the group consisting of subhuman primate antibody, murine monoclonal antibody, chimeric antibody, humanized antibody, and human antibody.

43. (Currently Amended) The method of claim 42, wherein said antibody is ~~the~~ **a** murine, chimeric, **human,** or humanized LL2 antibody.

44. (Canceled)

45. (Currently Amended) The method of claim 39, wherein said autoimmune disease is selected from the group consisting **of** acute idiopathic thrombocytopenic purpura, chronic idiopathic thrombocytopenic purpura, dermatomyositis, Sydenham's chorea, myasthenia gravis, systemic lupus erythematosus, lupus nephritis, rheumatic fever, polyglandular syndromes, bullous pemphigoid, diabetes mellitus, Henoch-Schonlein purpura, post-streptococcal nephritis, erythema nodosum, Takayasu's arteritis, Addison's disease, rheumatoid arthritis, multiple sclerosis, sarcoidosis, ulcerative colitis, erythema multiforme, IgA nephropathy, polyarteritis nodosa, ankylosing spondylitis, Goodpasture's syndrome, thromboangitis ubiterans, Sjogren's syndrome, primary biliary cirrhosis, Hashimoto's thyroiditis, thyrotoxicosis, scleroderma, chronic active hepatitis, polymyositis/dermatomyositis, polychondritis, pamphigus vulgaris, Wegener's granulomatosis, membranous nephropathy, amyotrophic lateral sclerosis, tabes dorsalis, giant cell arteritis/polymyalgia, pernicious anemia, rapidly progressive glomerulonephritis and fibrosing alveolitis.

46. (Previously Presented) The method of claim 39, further comprising separately administering a secondary therapeutic directed against T-cells, B-cells, plasma cells, or macrophages or inflammatory cytokines.

47. (Previously Presented) The method of claim 46, wherein said secondary therapeutic is administered prior to the administration of said therapeutic composition.

48. (Previously Presented) The method of claim 47, wherein said secondary therapeutic is administered concurrently with the administration of said therapeutic composition.

49. (Previously Presented) The method of claim 48, wherein said secondary therapeutic is administered after the administration of said therapeutic composition.

50. (Currently Amended) The method of claim 39, wherein said ~~antibody is~~ **therapeutic composition further comprises** an anti-CD20 antibody.

51. (Canceled).

52. (Previously presented) The method of claim 39, wherein said antibody is a naked antibody.

53. (Currently Amended) The method of claim 52, wherein said antibody is a ~~naked anti-CD22 antibody~~ **bispecific antibody**.

54-74. (Canceled).

75. (Currently Amended) ~~A~~ **The** method **according to claim 39,** ~~of treating multiple sclerosis, comprising administering to a subject with multiple sclerosis a~~ **wherein said** therapeutic composition ~~comprising~~ **comprises** a naked anti-CD20 antibody, a naked anti-CD22 antibody that binds with epitope B of the CD22 antigen, and a cytokine, wherein the two antibodies and the cytokine can be administered concurrently or in any order.

76. (Currently Amended) ~~A~~ **The** method according to 75, wherein the cytokine is IFN-ß.

77-106. (Canceled).

107. (New) The method according to claim 39, wherein said non-blocking anti-CD22 antibody binds a CD22 epitope selected from the group consisting of epitope A, epitope B, epitope C, epitope D and epitope E.

108. (New) The method according to claim 39, wherein said non-blocking anti-CD22 antibody binds the CD22 epitope recognized by the LL2 antibody.

109. (New) The method of claim 46, wherein said secondary therapeutic is selected from the group consisting of drugs, toxins, enzymes, hormones, cytokines, immunomodulators, boron compounds and therapeutic radioisotopes.

110. (New) The method of claim 39 wherein said autoimmune disease is selected from the group consisting of acute idiopathic thrombocytopenic purpura, chronic idiopathic thrombocytopenic purpura, myasthenia gravis, systemic lupus erythematosus, rheumatoid arthritis, multiple sclerosis, and Sjogren's syndrome.

111. (New) The method of claim 39, wherein said therapeutic composition comprises at least two monoclonal antibodies that bind with distinct CD22 epitopes, wherein one of said at least two monoclonal antibodies binds with a CD22 epitope selected from the group considering of epitope A, epitope B, epitope D, and epitope E and a second antibody binds with a different CD22 epitope selected from the group consisting of epitope A, epitope B, epitope C, epitope D and epitope E.